



ORIGINAL RESEARCH

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# Treatment results and prognostic factors of pediatric neuroblastoma: a retrospective study

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## Abstract

**Background:** We conducted a retrospective analysis to investigate treatment results and prognostic factors of pediatric neuroblastoma patients.

**Methods:** This retrospective study was carried out analyzing the medical records of patients with the pathological diagnosis of neuroblastoma seen at South Egypt Cancer Institute, Assiut University during the period from January 2001 and January 2010. After induction chemotherapy, response according to international neuroblastoma response criteria was assessed. Radiotherapy to patients with residual primary tumor was applied. Overall and event free survival (OAS and EFS) rates were estimated using Graphed prism program. The Log-rank test was used to examine differences in OAS and EFS rates. Cox-regression multivariate analysis was done to determine the independent prognostic factors affecting survival rates.

**Results:** Fifty three cases were analyzed. The median follow-up duration was 32 months and ranged from 2 to 84 months. The 3-year OAS and EFS rates were 39.4% and 29.3% respectively. Poor prognostic factors included age >1 year of age, N-MYC amplification, and high risk group. The majority of patients (68%) presented in high risk group, where treatment outcome was poor, as only 21% of patients survived for 3 year.

**Conclusion:** Multivariate analysis confirmed only the association between survival and risk group. However, in univariate analysis, local radiation therapy resulted in significant survival improvement. Therefore, radiotherapy should be given to patients with residual tumor evident after induction chemotherapy and surgery. Future attempts to improve OAS in high risk group patients with aggressive chemotherapy and bone marrow transplantation should be considered.

## Introduction

Neuroblastoma is the third most common malignancy in childhood and accounts for at least 15 percent of cancer-related deaths in children [1]. Despite concerted clinical and scientific efforts, prognoses of patients with high-risk neuroblastoma remain poor. About 30 to 35% of patients with high-risk disease who are older than 1 year survive more than 5 years [2,3].

Children with neuroblastoma exhibit marked variability in outcome when the disease is categorized by age, stage, and biologic characteristics [4,5]. Efforts to improve the outcome of patients with neuroblastoma have focused on identification of risk groups based on

clinical and biologic variables as well as intensification of therapy for high-risk cases [6].

## Patients and Methods

### Study subjects

This retrospective study was carried out analyzing the medical records of patients with the pathological diagnosis of neuroblastoma seen at the in the Pediatric Oncology, Clinical pathology and Radiotherapy Departments, SECI, Assiut University during the period from January 2001 and January 2010. Informed consent was obtained for all patients and the treatment decision was approved by institutional review board at our center. Eligible patients had histologically confirmed, neuroblastoma, and were previously untreated. For each patient, evaluation was done by history and examination, routine laboratory investigations, and imaging studies in the form of CT scan with contrast for the local disease, and bone scan.

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Bone marrow aspirate and biopsy was done for all patients. Histopathologic diagnosis was obtained from presenting mass and for uncertain cases, immunohistochemistry by neuron specific enolase was done. Patients were staged by international neuroblastoma staging system [7]. N-MYC amplification was done for all patients, using FISH in paraffin-embedded tissue sections.

#### Treatment Schedule

For patients with stage IVs; nine courses of doxorubicin (1 mg/kg) and cyclophosphamide (33 mg/kg) regimen (every 3 weeks), were given. For patients with stage III and IV disease; treatment included six courses of OPEC alternating with OJEC chemotherapy [table 1]. After the third and six courses, patients were evaluated for chemotherapy response by clinical examination, bone marrow aspirate and biopsy, CT scan of the local site and bone scan according to the initial presentation. Response according to international neuroblastoma response criteria [7] was assessed. Radiotherapy was given to patients with residual primary tumor, and for patients with cord compression as palliative treatment. CT planning was done for determination of target volume and critical structures. The radiation target volume included 1 cm margin around all sites of residual disease evident after induction chemotherapy and surgery. Radiotherapy was delivered using 6-MeV photon beams and antero-posterior/postero-anterior fields with customized blocks. A total dose of 24 Gy was given in 16 fractions [1.5 Gy per fraction]; 5 fractions per week. For younger children who needed anesthesia for immobilization, a total dose of 21 Gy was administered in hypofractionated schedule; 3 fractions [2.1 Gy per fraction] per week for 10 fractions. The clinical target volume had to be covered by the 95% isodose calculated for the reference point.

#### After-Therapy Monitoring

After completion of therapy, patients were followed up regularly every 3 months for 1 year, every 6 months for

the next 2 years, and annually thereafter. Follow-up examinations included physical examination, and routine laboratory studies. CT scans, and bone scan were done according to initial presentation. Assessment for treatment-related organ toxicity was done during treatment and after completion of therapy during follow up.

#### Statistical Methods

The study cutoff point was January 2010. Overall survival (OAS) was defined as the interval from enrollment to the date of death from any cause or to last follow-up. Event free survival (EFS) was defined as the interval from enrollment of patients to the date of relapse, progression, or death from any cause or to the date of last follow-up. Overall and event free survival rates were estimated using Graphed prism program. The Log-rank test was used to examine differences in OAS and EFS rates. Cox-regression multivariate analysis was done to determine the independent prognostic factors affecting survival rates.

#### Results

##### Patients' Characteristics [Table 2]

The median age of patients was 2 years (range: 3 months - 13 years). Twelve patients (22.6%) were under one year of age, and 41 patients (77.4%) above 1 year (65%). Thirty five patients were male (66%) and 18 were females (34%). The majority of patients were  $\geq 1$  year of age (41 patients; 77.4%), males (35 patients; 66%), presented with stages III & IV (49 patients; 92.5%), and with suprarenal tumor (46 patients; 86.8%). Low risk group ( $n = 4$ , 7.5%) included patients with stage IVs disease. Intermediate risk group ( $n = 13$ , 24.6%) included patients of  $<1$  year of age with III disease and non amplified N-MYC ( $n = 12$ ) as well as one patient of  $<1$  year of age with stage IV disease and non amplified N-MYC. High risk group ( $n = 36$ , 68%) included one patient of  $>1$  year of age with stage III and amplified N-MYC, one patient  $<1$  year of age with stage IV disease and amplified N-MYC and patients  $>1$  year of age

**Table 1 Chemotherapy protocols.**

OPEC regimen	Dose	Route and duration of administration	Timing
Vi Vincristine	1.5 mg/m <sup>2</sup>	i.v.	Day 1
Cisplatin	80 mg/m <sup>2</sup>	i.v. (6 hours infusion)	Day 2
Etoposide	200 mg/m <sup>2</sup>	i.v. (2 hours infusion)	Day 4
Cyclophosphamide	600 mg/m <sup>2</sup>	i.v. (2 hours infusion)	Day 1
Alternating every 3 weeks with OJEC regimen			
Vincristine	1.5 mg/m <sup>2</sup>	i.v.	Day 1
Carboplatin	500 mg/m <sup>2</sup>	i.v. (6 hour infusion)	Day 1
Etoposide	200 mg/m <sup>2</sup>	i.v. (2 hours infusion)	Day 1
Cyclophosphamide	600 mg/m <sup>2</sup>	i.v. (2 hours infusion)	Day 1

**Table 2 Patients' characteristics.**

Variable	No	%
<b>1. Age at diagnosis:</b>		
a) <1 year	12	22.6
b) ≥ 1 year	41	77.4
<b>2. Sex</b>		
a) Males	35	18
b) Females	66	34
<b>3. Stage</b>		
a) III	13	24.6
b) IV	36	67.9
c) IVS	4	7.5
<b>4. Risk group</b>		
a) Low Risk	4	7.5
b) Intermediate Risk	13	24.6
c) High Risk	36	67.9
<b>5. Primary site</b>		
a) Adrenal glands	46	86.8
b) Paraspinal	4	7.5
c) Pelvis	2	3.8
d) Mediastinal	1	1.9
<b>6. Metastatic sites</b>		
a) No metastasis	13	24.5
b) Bone marrow	18	34
c) Bone marrow and bone	14	26.4
d) Bone marrow and brain	2	3.8
e) Bone marrow and lung	1	1.9
f) Bone marrow, bone and liver	1	1.9
g) Bone marrow, lung and liver	1	1.9
h) Bone	1	1.9
i) Liver	1	1.9
j) Subcutaneous nodules	1	1.9
<b>7. MYCN amplification</b>		
a) yes	11	20.8
b) no	42	79.2
<b>8. Local Radiotherapy administration</b>		
a) yes	25	47.2
b) no	28	52.8
<b>TOTAL</b>	<b>53</b>	<b>100</b>

with stage IV disease (n = 34). Thirty six patients (67.9%) had BM infiltration, and 14 patients (26.4%) had positive bone scan. N-MYC amplification was detected in only 11 patients (20.8%). Adrenalectomy was done in 12 patients, adrenalectomy with nephrectomy in 2 patients, and pelvic mass excision in 2 patients. Radiotherapy to the site of residual primary tumor was given in 25 patients (47.2%), while palliative irradiation was given to 12 patients with painful bone metastases. The median follow-up from the date of enrollment was 32 months and ranged from 2 to 84 months.

#### For patients <1 year of age (n = 12)

##### For patients with stage IVs (n = 4)

After nine courses of doxorubicin and cyclophosphamide regimen, these patients achieved CR.

##### For patients with stage III (n = 6)

After 3 courses of OPEC/OJEC regimen, all 6 patients achieved PR. After 3 additional chemotherapy courses, 4 patients underwent surgical resection and the other 2 patients were irresectable. All 6 patients had received local radiotherapy.

##### For patients with stage IV (n = 2)

These 2 patients achieved PR after 6 courses of OPEC/OJEC regimen. They underwent surgical resection, but with residual disease. They had received local radiation therapy.

#### For patients ≥1 year of age (n = 41)

##### For patients with stage III (n = 7)

After 3 courses of OPEC/OJEC regimen, one patient died during chemotherapy at 2 months follow up, and 6 patients showed mixed response, but were irresectable. After additional 3 chemotherapy courses, they became resectable, underwent surgical resection, but with residual disease and received radiotherapy on residual tumor.

##### For patients with stage IV (n = 34)

After 3 courses of OPEC/OJEC regimen, 5 patients lost follow up, and 9 patients showed rapidly progressive disease with deteriorated general condition and eventually died; 2 of them died during chemotherapy and 7 shortly after it at follow up range of 4-8 months. For these 7 patients palliative radiotherapy was given to sites of painful bone metastases. Six patients had progressive disease and 14 showed mixed responses. After additional 3 chemotherapy courses, 6 patients with progressive disease showed stable disease, but were irresectable and received local radiotherapy. Four patients of those with mixed response showed stable disease and surgical resection was attempted. These 4 patients died in the early postoperative period (3 months). Ten patients of those with mixed response showed stable disease, but were irresectable. Five of these 10 patients showed deteriorated general condition, received palliative radiotherapy to sites of painful bone metastases, and eventually died (at a follow up range of 9-10 months). The remaining 5 patients were given local radiotherapy.

#### Survival analysis [Table 3, 4 and Figure 1, 2, 3 and 4]

Three-year OAS and EFS rates were 39.4% and 29.3% respectively, for all patients. Patients with <1 year of age showed 3-year OAS and EFS of 80% compared to 28.9% (p = 0.0005, HR: 0.26, 95% CI: 0.12-0.55) and 16.2% (p = 0.0004, HR: 0.26, 95% CI: 0.13-0.55) respectively, for older patients. For patients with low, intermediate and

**Table 3 Univariate analysis of prognostic factors affecting OAS and EFS rates.**

P value	3-year EFS	P value	3-year OAS	factor
<b>1-Age</b>		<b>P = 0.0005</b>		<b>P = 0.0004</b>
1 year (12)>	80%	HR:0.26,95%CI;	80%	HR:0.26,95%CI;
≥1 years (41)	28.9%	0.12-0.55	16.2%	0.12-0.55
<b>2-Sex</b>		<b>P = 0.902</b>		<b>P = 0.84</b>
Males (35)	41.4%	HR:1.05,95%CI;	30.9%	HR: 1.09, 95%CI;
Females (18)	34.7%	0.502-2.184	28.1%	0.52 - 2.24
<b>3-Risk group</b>		<b>p &lt; 0.0001</b>		<b>p &lt; 0.0001</b>
Low Risk (4)	100%		100%	
Intermediate Risk (13)	74.6%		74.6%	
High Risk (36)	20.7%		6.1%	
<b>4-Primary site</b>		<b>P = 0.059</b>		<b>P = 0.23</b>
a) Adrenal glands (46)	32.7%	HR:2.44,95%CI;	27%	HR: 1.77, 95%CI;
b)Other sites (7)	71.4%	0.97-6.16	57.1%	0.697 - 4.530
<b>5-MYCN amplification</b>		<b>P = 0.0027</b>		<b>P &lt; 0.0001</b>
yes (11)	8.3%	HR:4.58; 95%CI;	0	HR:10.64; 95%CI;
no (42)	49%	1.69 - 12.37	37.2%	3.47 - 82.63
<b>6-Local RT administration</b>		<b>p = 0.0078</b>		<b>p = 0.032</b>
Yes (25)	57.6%	HR:0.39; 95%CI;	42.5%	HR:0.47; 95%CI;
No (28)	23.4%	0.19 - 0.78	19.5%	0.24-0.94

high risk diseases, 3-year OAS rates were 100%, 74.6% and 20.7% respectively ( $p < 0.0001$ ) and 3-year EFS rates were 100%, 74.6% and 6.1% respectively ( $p < 0.0001$ ). Multivariate Cox regression analysis confirmed only the association between risk group and both OAS ( $p = < 0.0001$ , HR: 27.96, 95% CI: 5.59-139.87) and EFS ( $p = < 0.0001$ , HR: 38.08, 95% CI: 6.87-210.84) rates. Patients with N-MYC amplification showed 3-year OAS and EFS rates of 8.3% and 0% respectively compared to 49% ( $p = 0.0027$ , HR: 4.58, 95% CI: 1.69 - 12.37) and 37.2% ( $p < 0.0001$ , HR: 10.64, 95% CI: 3.47 - 32.63) respectively for those no MYCN amplification. For patients who were given local radiotherapy, 3-year OAS was 57.6% compared to 23.4% for those who were not given RT ( $p = 0.0078$ , HR: 0.39, 95% CI: 0.19-0.78) while the 3-year EFS rate was 42.5% for irradiated patients compared to 19.5% for non irradiated patients ( $p = 0.032$ , HR:0.47; 95%CI; 0.24-0.94).

#### Acute Toxicity and Late Effects [Table 5]

Therapy was well tolerated; where only two patients (of high risk group) died from septicemia due to febrile

neutropenia. There was also one patient (of high risk group) developed chest infection after relapse. Chest X ray, CT chest and pulmonary functions tests revealed pulmonary fibrosis and this patient died in relapse. There is no late toxicity developed during the study period. Regarding grade III hematological toxicity, leucopenia occurred in 24 patients (45.3%), and thrombocytopenia in 21 patients (39.6%). Regarding grade III non hematological toxicity; infection in 18 patients, increased liver enzyme in 18 patients (34%), ototoxicity in 7 patients (13.2%) and increased serum creatinine in 4 patients (7.6%).

#### Discussion

Most children with low risk neuroblastoma can be cured with surgery alone [8]. Most infants with disseminated disease have favorable outcomes following treatment with chemotherapy and surgery [4]. In contrast, the majority of children with high risk disease die from progressive disease despite intensive multimodality therapy. Current multimodality protocols for high-risk neuroblastoma patients have incorporated radiation to the primary disease site [9]. It was suggested that radiation to the primary tumor bed might benefit patients with gross residual disease at the time of radiation [10], with irradiation of active tumor residual instead of initial tumor extension [11].

In the current study, the median age of patients was 2 years and was confirmed by Papaioannou and McHugh [12] who stated that the most undifferentiated

**Table 4 Results of the multivariate Cox regression analysis.**

Factor	Significance	HR	95% CI
<b>OAS</b>			
Risk group	<0.0001	27.96	5.59 - 139.87
<b>PFS</b>			
Risk group	<0.0001	38.08	6.87 - 210.84

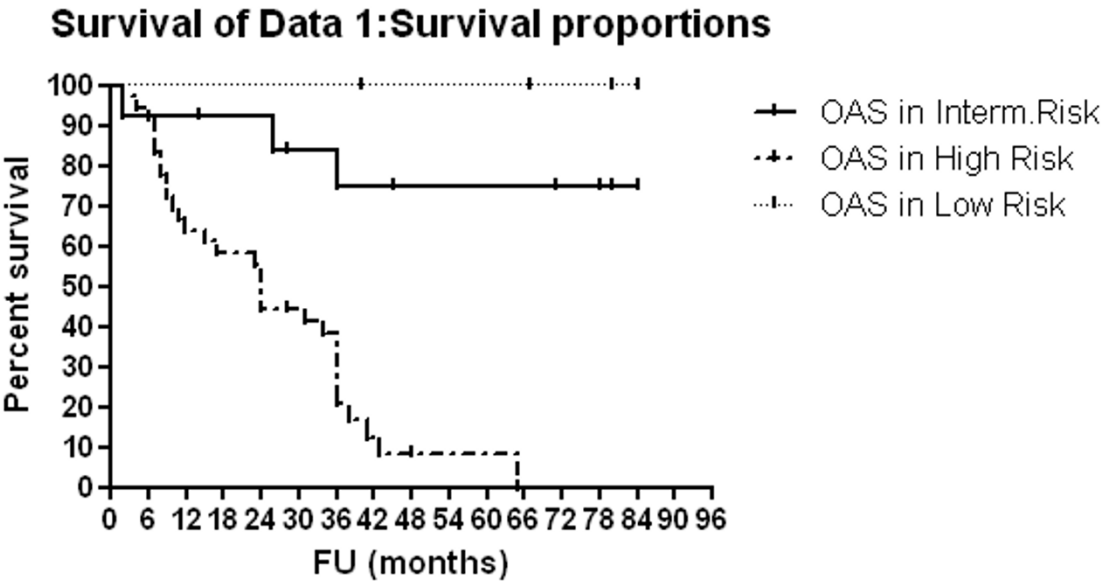


Figure 1 OAS according to risk group.

and aggressive neuroblastoma presents in young children with the median age  $\leq 2$  years. Compared with the current study, the reported studies, showed higher incidence of infants (45%) [13] and lower male to female ratio (1.3: 1) [14] and this may be explained by higher total number of patients (n = 500 - 594) in their studies.

Most of our patients presented with stage III & IV (92.5%). In the reported studies, it was found that the stages most frequently encountered at diagnosis were stages III and IV [15,16], and that 70% of the cases of children with NB presented with metastasis at diagnosis [17]. The distribution of our patients according to risk stratification (low risk in 7.5% and high risk in 68%) was different from that of Oberthuer et al. [18] (low risk in 50% and high risk in 40%) who used a gene expression-based classifier for neuroblastoma patients to reliably predicts courses of the disease. N-MYC amplification was detected in only 11 patients (21%) in the present study. This is lower than that found by Hass-Kogan et al., (27%) [10] and Schmid et al., (31%) [4], probably due

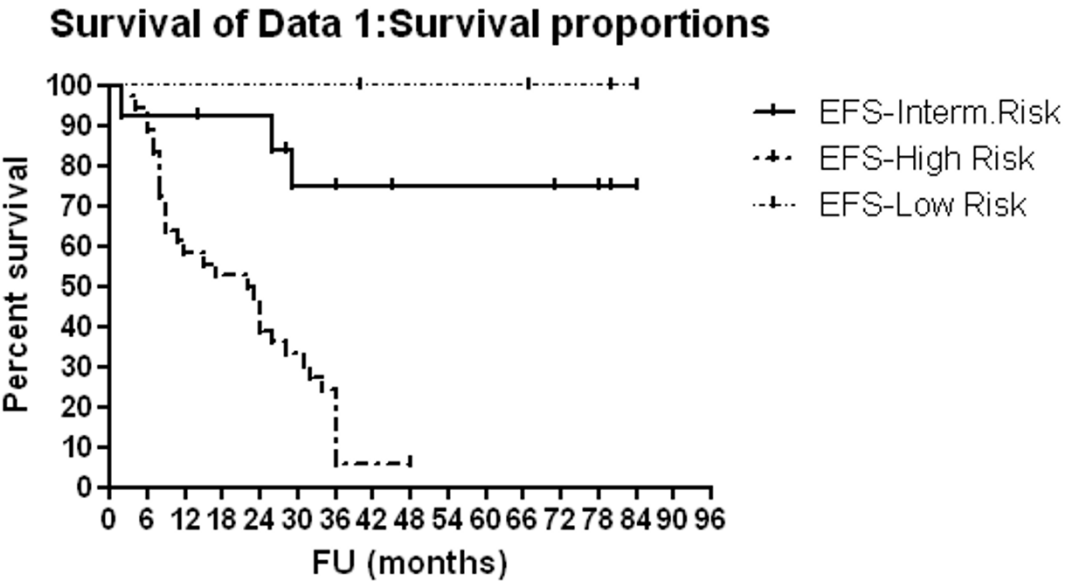
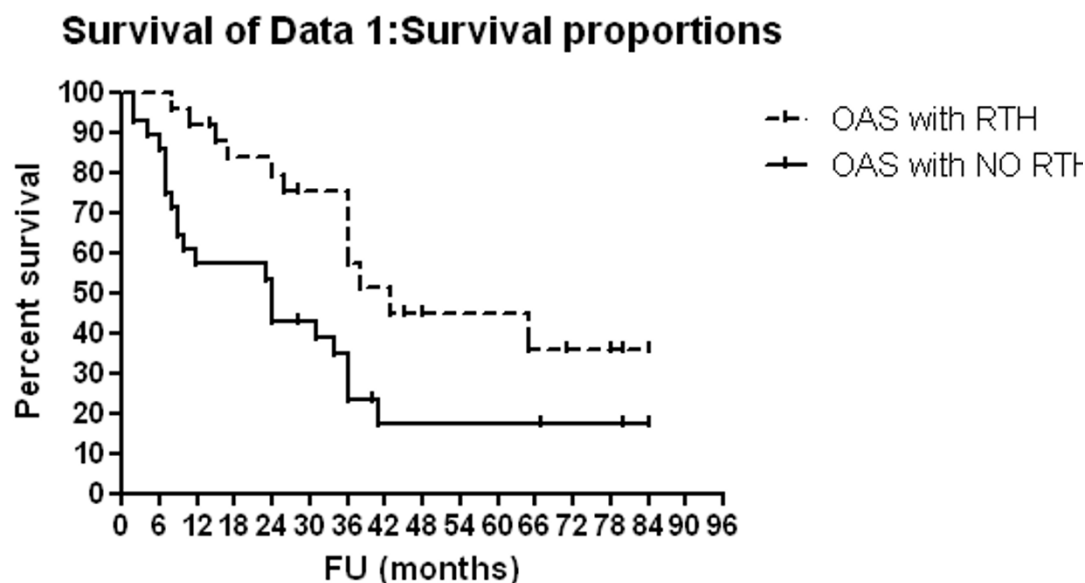


Figure 2 EFS according to risk group.



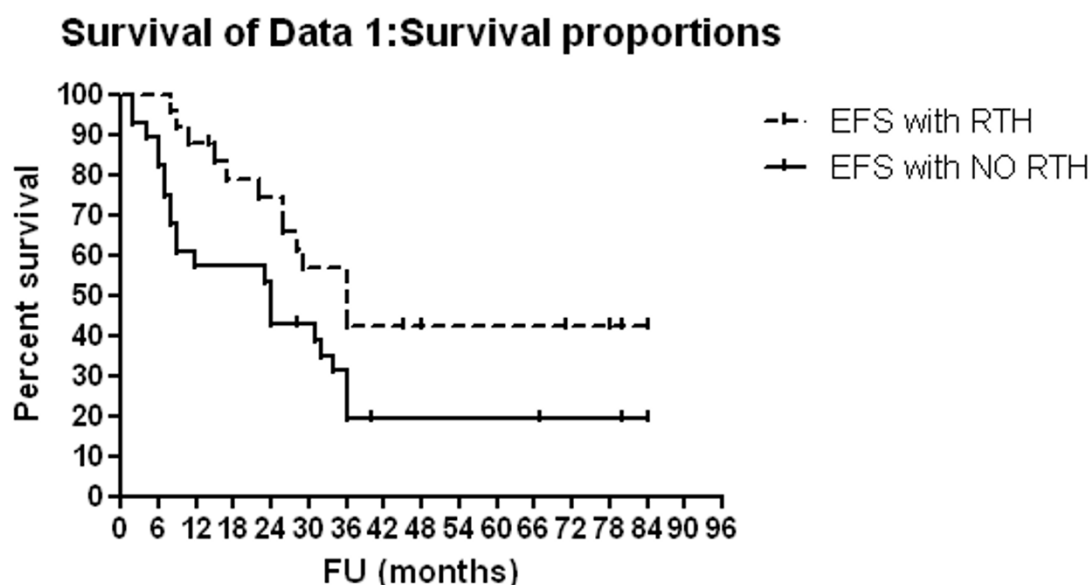
**Figure 3** OAS according to radiotherapy administration.

to higher total number of patients ( $n = 134 - 539$ ) in the reported studies [4,10].

The prognosis for neuroblastoma varies because of its peculiar biologic behavior [19]. The estimated OAS and EFS rates at 3 years for our patients were 39.4 and 29.3%, respectively. Three year OAS and EFS rates were significantly better in the 12 infants (80%) compared with the 41 children  $\geq 1$  year of age (29%;  $P = 0.0005$  and 16.2%;  $P = 0.0004$ , respectively). This was confirmed

by Evans and D'Angio, [20] who stated that, age younger than 1 year is a strongly favorable factor by itself. It was stated that OAS and EFS rates were significantly better in infants compared with children  $\geq 1$  year of age [21,22]. However, Schmidt et al [23], showed that EFS was 74% for the 12- to 18-month age group compared with 31% for those 18 to 24 months of age ( $P = .008$ ).

The COG stratified patients into low, intermediate, or high risk categories based on prognostic features.



**Figure 4** EFS according to radiotherapy administration.



**Table 5 Treatment toxicity rates.**

Toxicity	Grade III	
	No	(95% CI)%
Leucopenia	24	45.3 (31.88 - 58.68)
Thrombocytopenia	21	39.6 (26.45 - 52.79)
Infection	18	34 (21.21 - 46.71)
Increased liver enzymes	18	34 (21.21 - 46.71)
Nausea and vomiting	10	18.9 (8.34 - 29.4)
Ototoxicity	7	13.2 (4.09 - 22.33)
Increased serum creatinine	4	7.6 (0.44 - 14.66)

Following a meeting in 2005 to review data obtained for 11054 patients treated between 1974 and 2002, age, stage at diagnosis, and N-MYC status were selected for initial risk grouping [24]. In the present study, for patients with low, intermediate and high risk diseases, 3-year OAS rates were 100%, 74.6% and 20.7% respectively ( $p < 0.0001$ ) and 3-year EFS rates were 100%, 74.6% and 6.1% respectively ( $p < 0.0001$ ). Multivariate analysis confirmed only the association between survival and risk group. In low risk patients, 4-year EFS and OAS rates of  $81\% \pm 4\%$  and  $98\% \pm 1.5\%$ , respectively, were reported in previous CCG studies following treatment with surgery alone [8]. Our figures in low risk group (100%, 3-year OAS and EFS rates) were comparable to that reported by Bowman et al, [25] where the estimated 3-year OAS rate for patients with hyperdiploid tumors that lacked N-MYC amplification was 96%. These findings supported the reduction-in-therapy approach that was tested in the current COG low-risk study. The overall objective of the COG low-risk study was to preserve the excellent survival rate for patients with low-risk NB by using surgery as the primary treatment approach, thereby minimizing the risks of acute and long-term chemotherapy-related morbidity for the majority of these patients [26].

Survival rates in intermediate risk group (75%, 3-year OAS and EFS rates) in the present study was also comparable to that found in one of the reported study, where patients showed estimated 3-year EFS and OAS rates of 85% and 95% respectively [27].

Survival for high-risk children has improved modestly during the past 20 years, although cure rates remain low. Survival rates in high risk group (20.7%, 3-year OAS and 6% EFS rates) in the present study were very low. In the literatures, these patients had long-term survival of only 10 to 20 percent with combination chemotherapy, surgery, and local radiation therapy [1]. However, treatment approaches that used a combination of induction therapy, myeloablative consolidation therapy with stem cell support, and biologic therapy had improved 5-year survival rates from less than 15% to 40% [24].

In the present study, patients with N-MYC amplification showed statistically significant lower 3-year OAS and EFS rates (8.3% and 0%, respectively) than OAS (49%,  $p = 0.0027$ ) and EFS (37%,  $p < 0.0001$ ) in patients with no N-MYC amplification. This was consistent with reported series [26].

Regarding radiotherapy, outcome analysis focused on EFS and OAS rather than local control rates, since local treatment must aim to improve the survival of each individual patient [19]. The NB97 radiotherapy approach (36 Gy to residual tissue on the local tumor site) did compensate the outcome disadvantage of incomplete response to induction therapy. The authors discussed that radiotherapy might be able to compensate a suspected disadvantage of residual tumor [11]. Our data supported this hypothesis, as there were statistically significant higher 3-year OAS and EFS rates (57.6% and 42.5%, respectively) in patients who were given local radiotherapy than OAS (23%,  $p = 0.0078$ ) and EFS (19.5%,  $p = 0.032$ ) in patients with no local radiotherapy. This was matched with Simon et al., [11] where 3-year EFS and OAS rates were statistically significant lower if residual primary tumor was not irradiated than those in irradiated patients. This was also confirmed by Castleberry et al., [28] who used chemotherapy with or without radiotherapy. Overall survival (73% vs 41%) was better in the group receiving radiation. Therefore, it is recommended to give local radiotherapy to the residual primary tumor site [10].

Grade III leucopenia occurred in 24 patients (45.3%), and thrombocytopenia in 21 patients (39.6%). In the reported studies, grade 3/4 hematologic toxicity was more common (66 - 71%) and may be due to longer duration of induction chemotherapy [4] or more intensive chemotherapy course [27]. Infection was detected in 18 patients (34%) in the current study, which was within the range (24 - 58%) found in the reported studies [27,29]. Seven patients (13.2%) developed ototoxicity and 4 (7.6%) developed renal impairment in the present study. These figures were comparable to those found in CCG-3881, where renal and ototoxicity greater than grade 2 occurred in 3-6% of patients [4,29].

## Conclusion

The majority of patients (68%) presented in high risk group, where treatment outcome was poor. Multivariate analysis confirmed only the association between survival and risk group. However, in univariate analysis, there were statistically significant higher 3-year OAS and 3-year EFS rates in patients who were given local radiotherapy compared to survival rates in non irradiated patients. Therefore, radiotherapy should be given to patients with residual tumor evident after induction chemotherapy and surgery. Future attempts to improve

OAS in high risk group patients with aggressive chemotherapy and bone marrow transplantation should be considered.

## Conflict of interests

The authors declare that they have no competing interests.

## Acknowledgements

The authors received no financial or other support for the research reported in this manuscript

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## Authors' contributions

MIE carried out radiation therapy administration, follow up, statistical analysis, general coordination, drafting of the manuscript and writing of the final manuscript. AMA, and HAS carried out collection of patients' data, patient diagnosis, chemotherapy administration and follow up. EMZ identified N-MYC status using FISH technique. All authors have read and approved the final manuscript.

Received: 31 August 2010 Accepted: 24 December 2010

Published: 24 December 2010

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doi:10.1186/1755-7682-3-37

Cite this article as: El-Sayed et al.: Treatment results and prognostic factors of pediatric neuroblastoma: a retrospective study. *International Archives of Medicine* 2010 **3**:37.